

PATENT

Confirmation No.: 3346

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Application No.:

10/812,448

Stuyver

Applicant:

Filed:

March 29, 2004

TC/A.AU.:

1614

Examiner:

Unassigned

Docket No.:

08841.105059 PHA 2070 US

Customer No.:

20786

Title:

Compounds For the Treatment of Flaviviridae Infections

Mail Stop Amendment Commissioner for Patents P. O. Box 1450 Alexandria, VA 22313-1450

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Sir:

The citation of information on the accompanying Form PTO/SB/08A, "Information Disclosure Statement by Applicant" is made pursuant to 37 C.F.R. §§ 1.56, 1.97, and 1.98. A copy of each reference is enclosed. The citation of this information does not constitute an admission of priority or that any cited item is available as a reference, or a waiver of any right the applicant may have under applicable statutes, Rules of Practice in patent cases, or otherwise.

Applicant does not believe any fees are due because this paper is submitted before the mailing of a first Office action on the merits, as under 37 C.F.R. § 1.97(b)(3). However, the Commissioner is hereby authorized to charge any fees due or credit any overpayment, to Deposit Account No. 11-0980.

Respectfully submitted,

Madeline Johnston, Ph.D., Esq.

Reg. No. 36,174

Dated: November 5, 2004 King & Spalding LLP

191 Peachtree Street

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Docket No.: <u>08841.105059 PHA 2070 US</u>

Minikia D. Blair

Paralegal

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INFORMATION DISCLOSURE STATEMENT BY APPLICANT

(use as many sheets as necessary)

Sheet 1 of 6

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Application Number	10/812,448						
Filing Date	March 29, 2004						
First Named Inventor	Stuyver						
Group Art Unit	1614						
Examiner Name	Unassigned						
Attorney Docket Number	08841.105059 PHA 2070 US						

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			U	S. PATENT DOCUMENTS		
Examiner Initials *	Cite No. 1			Name of Patentee or Applicant of Cited Document	Date of Publication of Cited Document MM-DD-YYYY	Pages, Columns, Lines, Where Relevant Passages / Relevant Figures Appear
	AA	3,798,209	A	Witkowski et al.	03-19-1974	
	AB	Re. 29,835	Re.	Witkowski et al.	11-14-1978	
	AC.	4,211,771	A	Witkowski et al.	07-08-1980	
	AD	5,767,097	A	Tam	06-16-1998	
	AE	5,830,455	A	Valtuena et al.	11-03-1998	
	AF	6,063,628	A	Loeb et al.	05-16-2000	
	AG	6,063,772	A	Tam	05-26-2000	
	AH	6,150,337	A	Tam	11-21-2000	
	ΑI	6,348,587	B1	Schinazi et al.	02-19-2002	
	AJ	6,423,695	B1	Tam et al.	07-23-2002	
•	AK	6,455,508	B1	Ramasamy et al.	09-24-2002	
	AL	6,455,690	B1	Tam et al.	09-24-2002	
	AM	6,479,463	B1	Wang et al.	11-12-2002	
	AN	6,495,677	B1	Ramasamy et al.	12-17-2002	
	AO	6,509,320	B1	Wang et al.	01-21-2003	
	AP	6,573,248	B2	Ramasamy et al.	06-03-2003	
	AQ	6,660,721	B1	Devos et al.	12-09-2003	
	AR	6,777,395	B1	Bhat et al.	08-17-2004	
	AS	6,784,161	B1	Ismaili <i>et al</i> .	08-31-2004	
	AT	6,784,166	B1	Devos et al.	08-31-2004	
	AU	2002/0019363	A1	Ismaili <i>et al</i> .	02-14-2002	
	AV	2002/0055483	A1	Watanabe et al.	05-09-2002	
	AW	2002/0058635	A1	Averett	05-16-2002	
	AX	2002/0137696	A1	Tam	09-26-2002	
	AY	2002/0147160	A1	Bhat et al.	10-10-2002	
	ΑZ	2002/0147160	A1	Bhat et al.	10-10-2002	
	AZ	2002/0156030	A1	Ramasamy et al.	10-24-2002	
	AAA	2002/0198171	Al	Schinazi et al.	12-26-2002	
	AAB	2003/0008841	A1	Devos et al.	01-09-2003	
	AAC	2003/0028013	A1	Wang <i>et al</i> .	02-06-2003	
	AAD	2003/0050229	A1	Sommadossi et al.	03-13-2003	
	AAE	2003/0060400	A1	LaColla et al.	03-27-2003	
-	AAF	2003/0083307	A1	Devos et al.	05-01-2003	
	AAG	2003/0087873	A1	Stuyver et al.	05-08-2003	

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Submitted for	form 1449/PTO			Complete if Known		
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INI	FORMATION	DISCLO	DSURE	Filing Date	March 29, 2004	
	ATEMENT BY			First Named Inventor	Stuyver	
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Sheet	2	of	6	Attorney Docket Number	08841.105059 PHA 2070 US	

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			U	S. PATENT DOCUMENTS		
Examiner Initials *	Cite No. 1	U.S. Patent Docur Number K	nent ind Code ² (if known)	Name of Patentee or Applicant of Cited Document	Date of Publication of Cited Document MM-DD-YYYY	Pages, Columns, Lines, Where Relevant Passages / Relevant Figures Appear
	BA	2003/0119764	A1	Loeb et al.	06-26-2003	
	BB	2003/0124512	A1	Stuyver	07-03-2003	
	BC	2003/0225029	A1	Stuyver	12-04-2003	
	BD	2003/0236216	A1	Devos et al.	12-25-2003	
	BE	2004/0002476	A1	Stuyver et al.	01-01-2004	
	BF	2004/0002596	A1	Hong et al.	01-01-2004	
	BG	2004/0006002	A1	Sommadossi et al.	01-08-2004	
	BH	2004/0006007	A1	Gosselin et al.	01-08-2004	,
	BI	2004/0063622	A1	Sommadossi et al.	04-01-2004	
	BJ	2004/0067877	A1	Schinazi et al.	04-08-2004	
	BK	2004/0067901	A1	Bhat et al.	04-08-2004	
	BL	2004/0072788	A1	Bhat et al.	04-15-2004	
	BM	2004/0077587	A1	Sommadossi et al.	04-22-2004	·
	BN	2004/0082574	A1	Wang et al.	04-29-2004	
	ВО	2004/0097461	A1	Sommadossi et al.	05-20-2004	
	BP	2004/0097462	A1	Sommadossi et al.	05-20-2004	
	BQ	2004/0101535	A1	Sommadossi et al.	05-27-2004	
	BR	2004/0102414	A1	Sommadossi et al.	05-27-2004	
	BS	2004/0110717	A1	Carroll et al.	06-10-2004	
	BT	2004/0110718	A1	Devos et al.	06-20-2004	

	FOREIGN PATENT DOCUMENTS									
Examiner Initials *	Cite No. 1	For Office ³	Foreign Patent Document Office ³ Number Kind Code ² (if known)		Name of Patentee or Applicant of Cited Document	Date of Publication of Cited Document MM-DD-YYYY	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear	T _e		
	BU	JP	61-130299	Α	Daiken Koyo KK	06-18-1986				
	BV	wo	02/03997 🖊	A1	ICN Pharmaceuticals, Inc.	01-17-2001				
	BW	wo	02/048165 🖊	A2	Pharmasset Limited	06-20-2002				
	BX	wo	03/051899 🖊	A1	Ribapharm	06-26-2003				
	BY	wo	03/061385	A1	Ribapharm	07-31-2003				
	BZ	wo	03/061576	A2	Ribapharm	07-31-2003				
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	CA	wo	03/062256	A1	Ribapharm	07-31-2003						
	СВ	wo	03/062257	A1	Ribapharm	07-31-2003						
	CC	wo	2004/003138	´ A1	Merck & Co.; Isis Pharmaceuticals	01-08-2004						
	CD	wo	2004/007512	A2	Merck & Co.; Isis Pharmaceuticals	01-22-2004						
·	CE	wo	2004/009020	A2	Merck & Co.; Isis Pharmaceuticals	01-29-2004						
	CF	wo	2004/084796	A2	Pharmasset Limited	10-07-2004						

		OTHER PRIOR ART – NON PATENT LITERATURE DOCUMENTS				
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	CG	AGBARIA, R., et al., "Antiproliferative effects of cyclopentenyl cytosine (NSC 375575) in human glioblastoma cells," Oncol Res., 9:111-118 (1997).				
	СН	ALI, S.M., et al., "Efficient enantioseletive syntheses of carbocyclic nucleoside and prostaglandin synthons," <i>Tetrahedron Letters</i> , 31(11):1509-1512 (1990).				
	BALZARINI, J., et al., "Effect of antimetabolite drugs of nucleotide metabolism on the a immunodeficiency virus activity of nucleoside reverse transcriptase inhibitors," Pharmace 87:175-187 (2000).					
	CJ	BATTAGLIA, A.M., et al., "Combination therapy with interferon and ribavirin in the treatment of chronic hepatitis C infection," Ann. Pharmacother. 34(4):487-494 (April 2000).				
	СК	BERENGUER, M., et al. "Hepatitis C virus in the transplant setting," Antivir. Ther. 3(Suppl. 3):125-136 (1998).				
	CL	BIANCHI, V., et al., "Inhibition of ribonucleotide reductase by 2'-substituted deoxycytidine analogs: possible application in AIDS treatment," <i>Proc. Natl. Acad. Sci. U.S.A.</i> , 91:8403-8407 (1994).				
	СМ	BUDAVARI, S., Editor, <u>The Merck Index</u> , <u>11th edition</u> , Merck & Co., Inc., Rahway, NJ, p1304, 1989 [Ribavirin (1-β-D-ribofuranosyl-1-1,2,4-triazole-3-carboxamide) / Virazole TM].				
	CN	CARROLL, S.S., et al., "Inhibition of hepatitis C virus RNA replication by 2'-modified nucleoside analogs," J. Biol. Chem., 278(14):11979-11984 (April 4, 2003).				
	со	CHU, C.K., et al., "Nucleosides. 117. synthesis of 4-oxo-B-(β-D-ribofuranosyl)-3H-pyrazolo[1,5a]-1,3,5-triazine (OPTR) via 3-amino-2N-carbamoyl-4-(β-D-ribofuranosyl)pyrazole (ACPR) derivatives," J. Heterocycl. Chem., 17:1435-1439 (1980).				
	СР	CRANCE, J.M., et al., "Inhibition of sandfly fever Sicilian virus (Phlebovirus) replication in vitro by antiviral compounds," Res. Virol., 148:353-365 (1997).				

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IN	FORMATION I	DISCLO	OSURE	Filing Date	March 29, 2004
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				Examiner Name	Unassigned
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		OTHER PRIOR ART – NON PATENT LITERATURE DOCUMENTS	
Examiner Initials *	Cite No. 1	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	T ⁶
	DA	DAVIS, G.L., "Current therapy for chronic hepatitis C," Gastroenterology, 118(2 Suppl. 1):S104-S114 (2000).	
	DB	De CLERCQ, E., et al. "Broad-spectrum antiviral and cytocidal activity of cyclopentenylcytosine, a carbocyclic nucleoside targeted at CTP synthetase," <i>Biochem. Pharmacol.</i> , 41(12):1821-1829 (1991).	
	DC	DOLLINGER, M.R., et al., "Analogs of 1-β-D-arabinofuranosylcytosine. Studies of mechanisms of action in Burkit's cell culture and mouse leukemia, and in vitro deamination studies," Biochemical Pharmacology, 16(4):689-706 (April 1967).	
	DD	DUSCHINSKY, R., et al., "Nucleosides. XXXVII. 5,6-substituted 5-fluorodihydropyrimidines and their 2'-deoxyribonucleosides," J. Med. Chem., 10(1):47-58 (January 1967).	
	DE	HEINEMANN, V., et al., "Gemcitabine: A modulator of intracellular nucleotide and deoxynucleotide metabolism," Semin. Oncol., 22:11-18 (1995).	
	DF	HOSHINO, J., et al., "Suppression of nuclear ADP-ribosyltransferase activity in Ehrlich ascites tumor cells by 5-azacytidine and its analogs," Biochemical and Biophysical Research Communications, 142(2):468-474 (January 30 1987).	
	DG	IKEHARA, M., et al., "Studies of nucleosides and nucleotides. XXXII. Purine cyclonucleosides. 3. Synthesis of 2'-deoxy- and 3'-deoxyadenosine from adenosine," <i>Chem. Pharm. Bull.</i> , 15(1):94-100 (January 1967).	:
	DH	KABAT, M.M., et al., "Synthesis of 5-beta-D-ribofuranosylnicotinamide and its N-methyl derivative. The isosteric and isoelectronic analogues of nicotinamide nucleoside," J. Med. Chem., 30(5):924-927 (May 1987).	
	DI	KANEKO, M., et al., "Synthesis and properties of 8,2'-cyclothioguanosine and related compounds," <i>Chem. Pharm. Bull.</i> , 20(3):635-637 (1972).	
	DJ	KAZIMIERCZUK, Z., et al., "Synthesis of 2'-deoxytubercidin, 2'-deoxyadenosine, and related 2'-deoxynucleosides via a novel direct stereospecific sodium salt glycosylation procedure," J. Am. Chem. Soc., 106(21):6379-6382 (1984).	
	DK	LEE, W. W., et al., "Potential anticancer agents. LV. Synthesis of 3'-amino-2',3'-dideoxyadenosine and related analogs," J. Am. Chem. Soc., 83:1906-1911 (April 20, 1961).	
	DL	LIN, T. S., et al., "Synthesis and anticancer activity of various 3'-deoxy pyrimidine nucleoside analogs, and crystal structure of 1-(3-deoxy-β-D-threo-pentofuranosyl)cytosine," J. Med. Chem., 34(2):693-701 (1991).	
	DM	LOCKSHIN, A., et al., "Selective cytotoxicity of 5-hydroxyuridine for human colon adenocarcinoma cells," Cancer Treatment Reports, 69(7-8):845-850 (July-August 1985).	
	DN	LOHMANN, V., et al., "Biochemical and kinetic analyses of NS5B RNA-dependent RNA polymerase of the hepatitis C virus," Virology, 249(1):108-118 (September 15, 1998).	

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Examiner nitials *	Cite No. 1	sium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.							
	EA	MARKLAND, W., et al., "Broad-spectrum antiviral activity of the IMP dehydrogenase inhibitor VX-497: A comparison with ribavirin and demonstration of antiviral additivity with alpha interferon," Antimicrob. Agents Chemother., 44(4):859-866 (April 2000).							
	EB	MARUMOTO, R., et al., "One-step halogenation at the 2'-position of uridine, and related reactions of cytidine and N ⁴ -acetylcytidine," Chem. Pharm. Bull., 22(1):128-134 (1974).	Ī						
	EC	MENDEZ, E., et al., "Infectious bovine viral diarrhea virus (strain NADL) RNA from stable cDNA clones: a cellular insert determines NS3 production and viral cytopathogenicity," J. Virol., 72(6):4737-4745 (1998).							
	ED	MORREY, J.D., et al., "Identification of active antiviral compounds against a New York isolate of West Nile virus," Antiviral Res., 55:107-116 (2002).							
	EE	NEYTS, J., et al., "Use of the yellow fever virus vaccine strain 17D for the study of strategies for the treatment of yellow fever virus infections," <i>Antiviral Res.</i> , 30:125-132 (1996).							
	EF	NIEDBALLA, U., et al., "A general synthesis of N-glycosides. 6. On the mechanism of the stannic chloride catalyzed silyl Hilbert-Johnson reaction," J. Org. Chem., 41(12):2084-2086 (June 11, 1976).							
	EG	OZOLS, A. M., et al., "Aminonucleosides and the derivatives. VI. A new synthesis of 1,2,5-tri-I-O-3-azido-3-deoxy-β-D-ribofuranose," Synthesis, 1980:557-559 (1980).							
	ЕН	PANKIEWICZ, K. W., et al., "A synthesis of ψ -cytidine," Carbohydr. Res., 127(2):227-233 (April 15, 1984).							
	EI	PANKIEWICZ, K. W., et al., "Synthesis of 2,2'-anhydro-2-hydroxy- and 6,2'-anhydro-6-hydroxy-1-β-D-arabinofuranosylnicotinamide as conformationally restricted nicotinamide nucleoside analogs," <i>Nucleosides & Nucleotides</i> , 10:1333-1344 (1991).							
	EJ	PLUNKETT, W., et al., "Gemcitabine: metabolism, mechanisms of action, and self-potentiation," Semin. Oncol., 22:3-10 (1995).							
	EK	POLITI, P.M., et al., "Phase I clinical trial of continuous infusion cyclopentenyl cytosine," Cancer Chemother. Pharmacol., 36:513-523 (1995).	Ī						
	EL	ROBERTS, M., et al., "Uridine and cytidine derivatives," J. Am. Chem. Soc., 74:668-669 (February 5, 1952).							
	ЕМ	SERGUEEVA, Z. A., et al., "Rapid and selective reduction of amide group by borane-amine complexes in acyl protected nucleosides," <i>Nucleosides Nucleotides Nucleic Acids</i> , 19(1-2):275-282 (January-February 2000).							
	EN	SHI, J., et al., "Synthesis and biological evaluation of 2',3'-didehydro-2',3'-dideoxy-5-fluorocytidine (D4FC) analogues: Discovery of carbocyclic nucleoside triphosphates with potent inhibitory activity against HIV-1 reverse transcriptase," J. Med. Chem., 42(5):859-867 (1999).							
	EO	STAVBER, S., et al., "Room-temperature reactions of CsSO ₄ F with organic molecules containing heteroatoms," J. Chem. Soc. Chem. Commun., 1983:563-564 (1983).							

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	FA	STUYVER, L.J. et al., "Inhibitors of the IMPDH enzyme as potential anti-bovine viral diarrhea virus agents," Antiviral Chem. Chemother., 13:345-352 (2003).	
	FB	STUYVER, L.J., et al., "Ribonucleoside analogue that blocks the replication of bovine viral diarrhea and hepatitis C viruses in culture," Antimicrob. Agents Chemother., 47(1):244-254 (January 2003).	
	FC	SUHADOLNIK, R. J., et al., "Synthesis of 3'-deoxyuridine via transglycosylation of uracil with 3'-deoxyadenosine (cordycepin)," Carbohydr. Res., 61:545-548 (1978).	
	FD	SUTTLE, D.P., et al., "Coordinate overproduction of orotate phosphoribosyltransferase and orotidine-5'-phosphate decarboxylase in hamster cells resistant to pyrazofurin and 6-azauridine," J. Biol. Chem., 254:4602-4607 (1979).	
	FE	TANAKA, H., et al., "A simplified synthesis of 8-substituted purine nucleosides via lithiation of 6-chloro-9-(2,3-Q-isopropylidene-β-D-ribofuranosyl) purine," Chem. Pharm. Bull., 31(2):787-790 (1983).	
	FF	TANAKA, H., et al., "Regiospecific C-alkylation of uridine: A simple route to 6-alkyluridines," Tetrahedron Lett., 1979(49):4755-4758 (1979).	
:	FG	THURBER, T. C., et al., "A novel ring contraction of O ⁵ -6(S)-cyclo-5-diazouridine. Elimination of a ring carbonyl group in preference to diatomic nitrogen," J. Am. Chem. Soc., 95(9):3081-3082 (May 2, 1973).	
	FH	WACHSMAN, M., et al., "Anticytomegaloviral activity of methotrexate associated with preferential accumulation of drug by cytomegalovirus-infected cells," <i>Antimicrob. Agents Chemother.</i> , 40(2):433-436 (February 1996).	
	FI	WALKER, M.P., et al., "HCV RNA-dependent RNA polymerase as a target for antiviral development," Curr. Opin. Pharmacol., 2:1-7 (2002).	
	FJ	WALTON, E., et al., "3'-Deoxynucleosides. I. A synthesis of 3'-deoxyadenosine," J. Am. Chem. Soc., 86:2952 (1964).	
	FK	WATANABE, "The Chemistry of C-Nucleosides", in TOWNSEND, L. B., Ed., In "Chemistry of Nucleosides and Nucleotides", Plenum Publ., New York, Vol., 3, page 421-465 (1994).	
	FL	WOLFE, M.S., et al., "Enantiospecific syntheses of aristeromycin and neplanocin A," J. Org. Chem., 55:4712-4717 (1990).	

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